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(54) Porphyrin metal complex-albumin inclusion compound and oxygen carrier composition comprising the same

(57) A porphyrin metal complex-albumin inclusion compound wherein a substituted porphyrin metal complex comprising, as a central, coordinated metal, a transitional metal belonging to the fourth or fifth period of the periodic table, is included in the albumin, and an oxygen carrier composition comprising said inclusion compound as an active ingredient. The compound functions as an oxygen carrier which is superior in biocompatibility, capable of adsorption and desorption of oxygen under physiological conditions, and is rather easily produced at an industrial scale.

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(54) Porphyrin metal complex-albumin inclusion compound and oxygen carrier composition comprising the same

Porphyrinmetallkomplex-Albumin-Einschlussverbindung und diese enthaltender Sauerstoff-Träger

Composé d'inclusion d'un complexe de porphyrine-métaux et de l'albumine et porteur d'oxygène le contenant

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(56) References cited:
EP-A- 0 601 183 **WO-A-96/03426**

- **BONAVENTURA J ET AL: "OXYGEN-BINDING ALBUMINS: A NOVEL APPROACH TO BLOOD SUBSTITUTES" ARTIFICIAL CELLS, BLOOD SUBSTITUTES, AND IMMOBILIZATION BIOTECHNOLOGY, vol. 22, no. 5, 1 January 1994, page A83 XP000576501**
- **PATENT ABSTRACTS OF JAPAN vol. 9, no. 12 (C-261), 18 January 1985 & JP 59 162924 A (H. TSUCHIDA), 13 September 1984**
- **PATENT ABSTRACTS OF JAPAN vol. 18, no. 528 (C-1258), 6 October 1994 & JP 06 184156 A (RES INST FOR PROD DEV ET AL), 5 July 1994**
- **PATENT ABSTRACTS OF JAPAN vol. 18, no. 680 (C-1291), 21 December 1994 & JP 06 271577 A (RES INST OR PROD DEV), 27 September 1994**
- **J. P. COLLMAN ET AL: ""Picket Fence Porphyrins." Synthetic Models for Oxygen Binding Hemoproteins" JOURNAL OF THE AMERICAN CHEMICAL SOCIETY, vol. 97, January 1975 - March 1975, pages 1427-1439, XP002091695**
- **TRAYLOR T G ET AL: "SYNTHESES AND NMR CHARACTERIZATION OF CHELATED HEME MODELS OF HEMOPROTEINS" JOURNAL OF THE AMERICAN CHEMICAL SOCIETY, vol. 101, no. 22, 24 October 1979, pages 6716-6731, XP000575976**

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Description

[0001] The present invention relates to a novel inclusion compound comprising, as an oxygen adsorption-desorption site, a porphyrin metal complex included in an albumin, and to an oxygen carrier composition comprising said inclusion compound as an active ingredient.

[0002] A porphyrin iron (II) complex present in hemoglobin and myoglobin reversibly adsorbs and desorbs oxygen molecules. An attempt to impart an oxygen adsorption-desorption function similar to such natural porphyrin iron (II) complex by the use of a synthetic complex has been reported in a number of publications such as J.P. Collman, Accounts of Chemical Research, *10*, 265 (1977), F. Basolo, B.M. Hoffman, and J.A. Ibers, *ibid, 8*, 384 (1975). For the function of a synthetic porphyrin metal complex to be reproduced under physiological conditions (in physiological salt solution, pH 7.4, at room temperature or 37°C) and utilized in medical care (e.g., as oxygen supplying solution for artificial erythrocyte, organ preservation or oxygenator), the following requirements should be met. That is, (i) smallest possible concentration of imidazole derivatives widely used as axial base ligands for enhancing the oxygen binding capability, in view of the pharmacological action possessed by the imidazole derivatives, which sometimes causes high toxicity *in vivo*, and (ii) stable retention of oxygen coordinated complex by preventing the oxidation by the proton of the central metal and the oxidation via μ-oxo dimer as a result of not only making the porphyrin metal complex water-soluble, but also immobilization of the same in a microscopic hydrophobic environment.

[0003] With respect to the above-mentioned requirement (i), a porphyrin compound has already been synthesized having an imidazolyl group bonded to a porphyrin ring by a covalent bond, namely, a substituted porphyrin compound having an axial base in the molecule (resulting in the molar ratio of porphyrin/imidazole suppressed to the minimum necessary ratio of 1:1), and it has been clarified that this compound forms a stable oxygen coordinated complex (JP-A-271577/1994). When the axial base is bonded by an ester bond in this porphyrin complex, the compound exhibits high biodegradability, making itself highly advantageous for administration to living organisms. In other words, the development of a highly safe oxygen binding site (porphyrin metal complex) has been completed.

[0004] On the other hand, it is known that the above-mentioned requirement (ii) regarding the provision of a microscopic hydrophobic environment to the porphyrin metal complex can be met by utilizing a micelle containing a surfactant, or an endoplasmic reticulum having two molecular membranes of phospholipid. However, micelles are morphologically dynamic, inferior in stability as compared to an endoplasmic reticulum having two molecular membranes, and are capable of forming only a poorly hydrophobic environment. Therefore, an endoplasmic reticulum having two molecular membranes is frequently used as a carrier, which has a relatively stable shape and is capable of providing a sufficiently hydrophobic environment, so as to provide a hydrophobic environment to the complex. Thus, an oxygen carrier composition has been developed, which carries oxygen stably under physiological conditions as a result of dispersion, with high orientation, of porphyrin metal complex between the two molecular membranes of phospholipid endoplasmic reticulum.

[0005] Noting the probability of inclusion, with high orientation, of porphyrin metal complex in the hydrophobic environment of phospholipid having two molecular membranes, as achieved by the introduction of an alkyl substituent having hydrophilic ends onto the porphyrin ring, thereby to make an emphilic structure of the porphyrin, various emphilic porphyrin metal complexes have been synthesized and include-oriented in phospholipids having two molecular membranes, whereby a series of oxygen carrier compositions effective in aqueous phase systems have been developed (JP-A-101490/1984 and JP-A-213777/1983).

[0006] However, the use of a large amount of phospholipid to prepare such oxygen carrier composition leaves room for an improvement in terms of industrial scale production and biocompatibility inclusive of metabolism.

[0007] J. P. Bonaventura et al. report in an abstract published on page A83 in "Artificial Cells, Blood Substitutes, and Immobilization Biotechnology, vol. 22(5), 1994, about the testing of the hypothesis that a serum albumin containing a transition metal-based oxygen binding site can be prepared. It is mentioned that so-called "picket fence porphyrins" have been made.

[0008] EP-A-601 183 deals with an anti-HIV drug comprising at least one porphyrin derivative as the active ingredient.

[0009] JP-A-59162924 and JP-A-06271577 refer to a specific porphyrin derivative, however, there is nothing disclosed about the stabilization of these compounds by forming a complex with albumin.

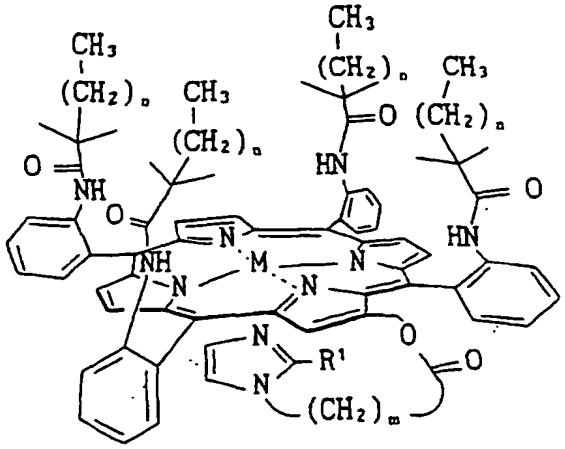
[0010] It is therefore an object of the present invention to provide a novel compound comprising, as an oxygen absorption-desorption site, a porphyrin metal complex, which is superior in biocompatibility and permits rather easy production at an industrial scale, and an oxygen carrier composition comprising the same.

[0011] This object could be achieved as a consequence of investigations of the molecular design of an oxygen carrier capable of stably carrying oxygen under physiological conditions, expression of the function thereof and high biocompatibility of a carrier capable of providing a hydrophobic environment to the porphyrin metal complex, wherein it was found that albumin which occupies 50-55% of plasma protein and which carries various compounds in the living body can provide a superior hydrophobic environment to the porphyrin metal complex, and that a porphyrin metal complex-albumin inclusion compound obtained by including a certain porphyrin metal complex in the albumin forms a stable

oxygen coordinated complex in water, and thus is able to function as an oxygen carrier.

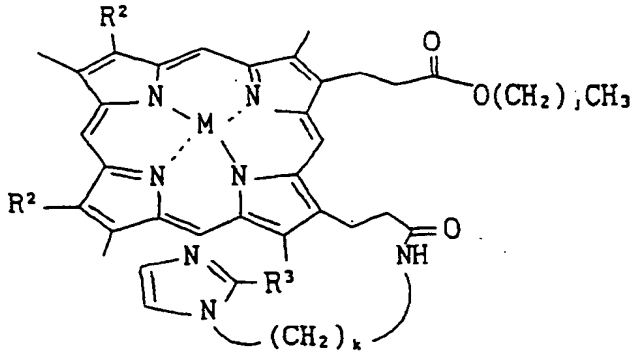
[0012] Accordingly, the present invention provides a porphyrin metal complex-albumin inclusion compound wherein a substituted porphyrin metal complex having a central coordinated metal is included in albumin.

[0013] The substituted porphyrin metal complex is a 5,10,15,20-tetra[α,α,α,α-o-(substituted amide)phenyl]porphyrin metal complex having the following formula (I):



(I)

wherein R^1 is hydrogen or methyl, M is an iron or cobalt ion, n is an integer of from 0 to 17 and m is an integer of from 3 to 17, or a porphyrin metal complex of the formula (II)



(II)

wherein the two R^2 groups are the same or different and each is a vinyl or a 1-alkyloxyethyl group of the formula : $-(CH_3)CHO(CH_2)_hCH_3$ wherein h is an integer of from 0 to 17, R^3 is hydrogen or methyl, M is an iron or cobalt ion, j is an integer of from 0 to 17 and k is an integer of from 3 to 10.

[0014] The porphyrin metal complex of the formula (I) or (II) which is an oxygen carrier comprises iron or cobalt as the central divalent metal M.

[0015] The origin of the albumin is not particularly limited, but it is preferably human serum albumin or recombinant human serum albumin. In particular, the recent progress in gene recombination has enabled the provision of a highly pure recombinant human serum albumin having the completely same structure; composition and physicochemical characteristics with human serum albumin [see Yokoyama and Ohmura, Clinical Molecular Medicine, *1*, 939 (1993)], and a porphyrin metal complex-recombinant human serum albumin inclusion compound wherein a substituted porphyrin metal complex has been included in the recombinant human serum albumin can be entirely prepared by synthesis, thus making industrial production easier.

[0016] In another aspect of the present invention, an oxygen carrier composition comprising, as an active ingredient, the above-mentioned porphyrin metal complex-albumin inclusion compound is provided.

[0017] The present invention is explained in more detail in the following.

[0018] In the present invention, the albumin including the porphyrin metal complex to be detailedly described later is a simple protein with its main function being control of colloidal osmotic pressure in blood, and also functions as a carrier protein of e.g. nutritive substances, metabolites thereof and drugs. The present invention relies on such non-specific binding capacity of albumin, which is absent in other proteins.

[0019] In addition, albumin is markedly advantageous in the application to the living body, particularly as an erythrocyte substitute, as compared to the system using phospholipid endoplasmic reticulum, since it is a blood plasma protein.

[0020] The origin of the albumin to be used in the present invention is not particularly limited and the albumin is exemplified by human serum albumin, recombinant human serum albumin and bovine serum albumin. In consideration of the application to humans, however, the use of human serum albumin or recombinant human serum albumin is desirable.

[0021] The substituted porphyrin metal complex to be included in albumin in the present invention is not subject to any particular limitation as long as it is a substituted porphyrin metal complex having, as a central coordinated metal iron or cobalt ion M an The central coordinated metal is preferably iron. An imidazole-bound substituted porphyrin metal complex is particularly preferable.

[0022] In a preferred embodiment of the present invention, the substituted porphyrin metal complex is expressed by the above-mentioned formula (I) or (II).

[0023] 5,10,15,20-tetrakis(o-pivalamidophenyl)porphyrinato copper (II) was prepared according to Collman et al., *J. Am. Chem. Soc.*, 1975, *97*, 1427.

[0024] Phosphorus oxychloride (3.5 ml, 37.3 mmol) was dropped into N,N-dimethylformamide (DMF) (3.5 ml) in an ice-water bath and the mixture was stirred for 1 hour at room temperature, giving a Vilsmeier reagent. To the porphyrin (0.2 g, 0.19 mmol) solution dissolved in dry CH_2Cl_2 (40 ml) was added dropwise at 25°C and refluxed for 15 hours, yielding a green solution. Saturated aqueous sodium acetate (50 ml) was added to the mixture at 25°C and further stirred for 3 hours at 40°C. The mixture was extracted by $CHCl_3$ and washed with water. After drying (Na_2SO_4), the organic layer was chromatographed on a silica gel flash column using $CHCl_3$-ethylacetate (6/1 v/v) as the eluent. The residue was dried at room temperature in vacuo to give a purple product, (2-formyl-5,10,15,20-tetrakis(o-pivalamidophenyl)porphyrinato copper (II), 0.16 g, 78%).

[0025] To a CH_2Cl_2 solution (20 ml) of (2-formyl-5,10,15,20-tetrakis(o-pivalamidophenyl)porphyrinato copper (II) (0.16 g), conc. H_2SO_4 (20 ml) was added and vigorously stirred for 10 min. The resulting green solution was added dropwise to CH_2Cl_2-ice-water (1/3 v/v) (800 ml) and neutralized by $NaHCO_3$ slowly. The mixture was extracted by $CHCl_3$ and washed with water. After drying (Na_2SO_4), the organic layer was chromatographed on a silica gel flash column using $CHCl_3$-ethylacetate (4/1 v/v) as the eluent. The residue was dried at room temperature in vacuo to give a purple product, (2-formyl-5,10,15,20-tetrakis(o-pivalamidophenyl)porphyrin, 0.12 g, 85%).

[0026] $NaBH_4$ (41.5 mg, 1.1 mmol) was added to a solution of CH_2Cl_2-MeOH (1/3 v/v) (8 ml) containing 2-formyl-5,10,15,20-tetrakis(o-pivalamidophenyl)porphyrin (0.11 g, 0.11 mmol) under argon and stirred for 15 min. After adding water to the solution, the mixture was extracted by $CHCl_3$ and washed with water. After drying (Na_2SO_4), the organic layer was chromatographed on a silica gel flash column using $CHCl_3$-MeOH (10/1 v/v) as the eluent. The residue was dried at room temperature in vacuo to give a purple product, (2-hydroxymethyl-5,10,15,20-tetrakis(o-pivalamidophenyl) porphyrin, 0.11 g, 94%).

[0027] 8-Imidazol-1-yloctanoic acid hydrochloride (90.1 mg, 0.37 mmol) and triethylamine (TEA, 0.1 ml, 0.74 mmol) were dissolved in dry DMF (4 ml) and stirred for 10 min. After removing TEA under reduced pressure, 2-hydroxymethyl-5,10,15,20-tetrakis(o-pivalamidophenyl)-porphyrin 76 mg, 73 μmol), 4-(N,N-dimethylamino)pyridine (DMAP, 4.5 mg, 37 μmol) and dicyclohexylcarbodiimide (DCC, 75.4 mg, 0.37 mmol) were added to the solution and stirred for 84 hours at room temperature in darkness. The mixture was chromatographed on a silica gel flash column using $CHCl_3$-MeOH (30/1 v/v) as the eluent. The residue was dried at room temperature in vacuo to give a purple product, 2-(8-imidazol-1-yloctanoyloxymethyl)-5,10,15,20-tetrakis(o-pivalamidophenyl)porphyrin (80 mg, 89%).

[0028] A dry tetrahydrofuran (THF, 50 ml) solution of 2-(8-imidazol-1-yloctanoyloxymethyl)-5,10,15,20-tetrakis(o-pivalamidophenyl)porphyrin (80 mg, 64 μmol) was added dropwise to anhydrous iron (II) bromide (0.86 g, 4 mmol) under dry argon and the mixture was refluxed under argon for 4 hours. The mixture was extracted by $CHCl_3$ and washed with water. After drying (Na_2SO_4), the organic layer was chromatographed on a silica gel flash column using $CHCl_3$-MeOH (4/1 v/v) as the eluent. The residue was dried at room temperature in vacuo to give a purple product, 2-(8-imidazol-1-yloctanoyloxymethyl)-5,10,15,20-tetrakis(o-pivalamidophenyl)porphyrinato iron (II) (100 mg, 84%).

[0029] The 5,10,15,20-tetra[α,α,α,α-o-(substituted amide)phenyl]-porphyrin metal complex of the formula (I) importantly comprises an alkyl imidazolyl group bonded to the 2-position of a porphyrin ring.

[0030] It was found that the porphyrin complex without such imidazolyl group in the molecule immediately degrades without forming a stable oxygen complex in an aqueous phase system, despite the addition of an excess external imidazole derivative (e.g., 1-methylimidazole).

[0031] The porphyrin of the formula (II) is a protoporphyrin IX derivative similar to hemoglobin in the living body, and

is superior in biocompatibility. Of the compounds of the formula (II), a porphyrin derivative wherein R^2 is vinyl can be synthesized by the method of T.G. Traylor et al., J. Am. Chem. Soc., *101*, 6716 (1979). A porphyrin derivative wherein R^2 is 1-alkyloxyethyl can be synthesized by, for example, the following process.

[0032] That is, 8,13-bis(1'-alkyloxyethyl)-3,7,12,17-tetramethyl-2,18-bis(2'-alkyloxycarbonylethyl)-21H,23H-porphyrin synthesized by the method described in Tsuchida et al., Chem. Lett., 1953, (1994) is dissolved in 1-5N hydrochloric acid, and the mixture is stirred for 1-24 hours at room temperature. When the porphyrin is insoluble in hydrochloric acid, an organic solvent such as tetrahydrofuran and acetone is added as necessary to homogeneously dissolve the porphyrin. After the completion of the reaction, the solvent is distilled away under reduced pressure, and the residue is extracted with e.g. chloroform, dichloromethane or benzene, which is followed by washing with pure water. Then, the organic layer is evaporated to dryness and subjected to separation and purification by silica gel column chromatography to give a monoester compound wherein one of the ester bonds has been hydrolyzed. This monoester compound is dissolved in anhydrous tetrahydrofuran, dimethylformamide and the like containing a base such as triethylamine, pyridine and 4-dimethylaminopyridine under a nitrogen atmosphere. Chloride pivalate is added at -20 - 30°C, preferably -20 - 0°C, and the mixture is stirred for 10-60 minutes. 1-(3-Aminoalkyl)imidazole is dropwise added and the mixture is reacted at -20-30°C for 1-24 hours. The solvent is removed under reduced pressure, and the residue is extracted with an organic solvent such as chloroform, dichloromethane and benzene, which is followed by washing with pure water. Then, the organic layer is evaporated to dryness under reduced pressure and subjected to separation and purification by silica gel column chromatography to give the desired 8,13-bis(1'-alkyloxyethyl)-2-(2'-alkyloxycarbonylethyl-18-(2'-(3"-imidazolyl)alkyl)-aminocarbonylethyl)-3,7,12,17-tetramethyl-21H,23H-porphyrin.

[0033] The introduction of the central metal, namely, the introduction of e.g. iron or cobalt is carried out by a conventional method [e.g., D. Dolphin ed., The porphyrin, Academic Press (1978)]. In general, a porphyrinato iron (III) complex is obtained from an iron complex and a porphyrinato cobalt (II) complex is obtained from a cobalt complex.

[0034] The substituted porphyrin metal complex-albumin inclusion compound of the present invention is prepared by the following step. That is, a porphyrin metal [e.g, 2-(8'-(N-imidazolyl)octanoyloxymethyl)-meso-tetra(α,α,α,α-o-pivalamidophenyl)porphyrinato iron (III) complex] is dissolved in a water-soluble solvent (e.g., dimethylformamide, dimethylsulfoxide and methanol) and an aqueous solution of albumin (e.g., human serum albumin) in, for example, water, ethanol, phosphate buffer (pH 5-9), physiological saline or Krebs-Ringer solution is added, followed by gentle shaking. The obtained aqueous dispersion is concentrated by ultrafiltration using, for example, an ultrafiltration membrane having a cut off molecular weight of 20,000-40,000, to about 10% of the total amount. Water is added and the ultrafiltration is repeated to give a porphyrin metal complex-albumin inclusion compound. This dispersion is free of sedimentation or coagulation even after preservation at 4-35°C for several months, and is stable.

[0035] When the central metal of the porphyrin metal complex is iron (III), an oxygen-binding activity can be imparted thereto by a conventional method such as addition of a reducing agent (e.g., aqueous solution of sodium dithionite or ascorbic acid) under a nitrogen atmosphere to reduce the central metal iron from trivalent to divalent.

[0036] Such reduction can be carried out by the addition of not only a reducing agent but also a palladium-carbon/hydrogen gas. For example, a porphyrin iron (III) complex is dissolved in dry e.g. dichloromethane, benzene or toluene; a small amount of palladium-carbon is added; and a hydrogen gas is sufficiently blown in at room temperature, whereby the central metal iron is reduced. After the reduction, the palladium-carbon is filtered off and the filtrate is dried in vacuo to give a porphyrin iron (II) compound.

[0037] The above-mentioned reduction can be carried out before inclusion reaction.

[0038] The porphyrin metal complex-albumin inclusion compound of the present invention thus obtained comprises a porphyrin metal complex included and immobilized in the inner hydrophobic region formed by albumin. The number of the porphyrin metal complex included and bonded to one mole of albumin can be determined by forming a Scatchard plot [C.J. Halfman, T. Nishida, Biochemistry, *11*, 3493 (1972)]. For example, the number of bonding of a substituted porphyrin metal complex (e.g., 2-(8'-(2"-methyl-1-imidazolyl))-octanoyloxymethyl-meso-tetra(α,α,α,α-o-pivalamidophenyl)-porphyrinato iron (II) complex) to the albumin (e.g., human serum albumin) is 1-3.

[0039] As has been stated, the oxidation by the proton of the central metal and oxidative degradation via μ-oxo dimer can be completely inhibited by the oxygen adsorption site (porphyrin metal complex) which has been immobilized in the inner hydrophobic environment of albumin. Consequently, the inclusion compound of the present invention can retain the stable oxygen coordinated complex in an aqueous phase system.

[0040] The porphyrin metal complex-albumin inclusion compound of the present invention comprises a synthetic substituted porphyrin metal complex as the oxygen binding site. Hence, its affinity for oxygen, toxicity, biodegradability and the like can be optionally controlled by varying the chemical structure of the substituted porphyrin metal complex. It has been clarified by the present invention that, while the inclusion in albumin requires a certain balance between hydrophilicity and hydrophobicity (polarity) and certain molecular volume of the porphyrin metal complex, a bulky molecule can be included and sufficiently exert the desired function, such as tetraphenylporphyrin derivative of the formula (I) having a molecular volume of 10-21 nm^3 which is about 2-10 times greater than the molecular volume of 2-3.5 nm^3 of protoporphyrin of the formula (II). This finding is expected to offer a new aspect of the molecular design of the albumin

complex.

[0041] As is evident from the foregoing explanation, the porphyrin metal complex-albumin inclusion compound of the present invention quickly forms a stable oxygen coordination complex upon contact with oxygen. This oxygen coordination complex is capable of adsorbing or desorbing oxygen according to the oxygen partial pressure. Such oxygen adsorption and desorption can be stably repeated reversibly according to the oxygen partial pressure. The oxygen bond dissociation is completed quickly, and the inclusion compound of the present invention can function as a semi-artificial oxygen carrier or entirely synthesized oxygen carrier when human serum albumin is used, which is capable of carrying oxygen in the blood streams in the living body.

[0042] According to a measurement, the half-life ($\tau_{1/2}$) of the oxygen coordination complex in a porphyrin iron (II) complex-albumin inclusion compound [e.g., 2-(8'-(2"-methyl-1-imidazolyl))octanoyloxymethyl-meso-tetra(α,α,α,α-o-pivalamidophenyl)porphyrinato iron (II) complex-human serum albumin inclusion compound] at 25°C was not less than 16 hours. In contrast, that of the oxygen coordination complex included between the two molecular layers of phospholipid endoplasmic reticulum under the same conditions was not less than 4 hours, and that in a micelle solution in which dispersion was effected by the use of surfactant Triton X-100 (trademark) was short and not more than one minute. Thus, the life time of an oxygen coordination complex can be greatly prolonged by the inclusion of the porphyrin metal complex in the inner hydrophobic environment of albumin.

[0043] The inclusion of the substituted porphyrin metal complex in an inner hydrophobic environment of albumin has led to the superior function of the porphyrin metal complex-albumin inclusion compound of the present invention as an oxygen carrier, as mentioned above. When the central metal is a transition metal belonging to the fourth or fifth period of the periodic table, moreover, the compound of the present invention can act as a catalyst in a wide range of reactions such as oxidative reduction, oxidative oxidation, active oxygen decomposition, optical activation of oxygen and oxygenation. In other words, the inclusion compound of the present invention is useful *per se* as a synthesized oxygen carrier, as well as a gas adsorbent, an oxygen adsorbent and desorbent, and a catalyst in oxidative oxidation or reduction, oxygenation and optical activation of oxygen. The inclusion compound of the present invention is easy to produce and suitable for industrial production when a recombinant human serum albumin is used.

[0044] The present invention is explained in the following by illustrative Examples to which the invention is not limited.

Example 1

[0045] According to the method described in JP-A- 271577/1994, 2-(8'-(2"-methyl-1"-imidazolyl)-octanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-pivalamidophenyl)porphyrinato iron (III) complex was obtained. To a solution of 0.14 mmol of this complex in dimethyl sulfoxide (12 ml) was added an aqueous solution of human serum albumin (0.1 mmol) in phosphate buffer (100 ml, pH 7.4, 1/30 mM), and the mixture was shaken. The mixture was concentrated to 20 ml by ultrafiltration (ultrafilter manufactured by ADVANTEC; cut off molecular weight 20,000). An aqueous solution of phosphate buffer (92 ml, pH 7.4, 1/30 mM) was added and the mixture was shaken, which was followed by concentration by ultrafiltration and dilution with an aqueous solution of phosphate buffer, whereby the desired dispersion of porphyrin iron (III) complex-albumin inclusion compound was obtained. This dispersion was preserved at room temperature or 4°C for several months. As a result, no sedimentation or coagulation was found, and the dispersion was stable.

[0046] A small amount of an aqueous solution of sodium dithionite was added to this dispersion under nitrogen atmosphere to reduce the central metal iron of the porphyrin complex to a divalent iron, whereby the objective dispersion of 2-(8'-(2"-methyl-1"-imidazolyl)-octanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-pivalamidophenyl)porphyrinato iron (II) complex-albumin inclusion compound was obtained.

[0047] A carbon monoxide gas was passed through this dispersion, and the light (500 W) was irradiated in an ice bath with nitrogen gas aeration to give a deoxy type complex. This dispersion was diluted 1/50-fold and transferred into a quartz spectrometry cell, and the cell was sealed under a nitrogen atmosphere. The visible absorption spectra thereof were λ_{max} : 563 nm, 542 nm, 439 nm, which correspond to the spectrum of a penta-coordinated deoxy type complex having one intramolecular base.

[0048] The spectrum immediately changed when an oxygen gas was passed through this dispersion, giving spectra of λ_{max} : 548 nm, 424 nm, which apparently suggests that the porphyrin iron (II) complex included in albumin had changed to an oxygenated complex. A nitrogen gas was passed through the dispersion of this oxygenated complex, as a result of which the visible absorption spectrum changed from the oxygen type spectrum to the deoxy type spectrum, thus confirming the reversible adsorption and desorption of oxygen. Repeats of alternative aeration of oxygen and nitrogen enabled repetitive adsorption and desorption of oxygen. The half life time of the above-mentioned oxygenated complex was not less than 16 hours at 25°C.

Example 2

[0049] In the same manner as in Example 1 except that 2-(18'-(N-imidazolyl)octadecanoyloxymethyl)-5,10,15,20-tet-

rakis(α,α,α,α-o-(2,2-dimethyloctadecanamidophenyl)porphyrinato iron (III) complex (1.4 mmol) was used instead of the porphyrin iron (III) complex used in Example 1 and bovine serum albumin was used instead of human serum albumin, a dispersion of 2-(18'-(N-imidazolyl)octadecanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-(2,2-dimethyloctadecanamidophenyl) porphyrinato iron (II) complex-albumin inclusion compound was prepared. The visible absorption spectra were λ_{max} : 562 nm, 541 nm, 436 nm, which correspond to the spectrum of a deoxy type complex. The aeration of oxygen gas in the same manner as in Example 1 provided visible absorption spectra (λ_{max} 544 nm, 423 nm) corresponding to the spectrum of an oxygenated complex. A nitrogen gas was passed through the dispersion of this oxygenated complex, whereby the original deoxy type spectrum was obtained, thus confirming the reversible adsorption and desorption of oxygen. The half life time of the above-mentioned oxygenated complex was not less than 15 hours at 25°C.

Example 3

[0050] 2-(8'-(N-Imidazolyl)octanoyloxymethyl)-5,10,15,20-tetrakis-(α,α,α,α-o-pivalamidophenyl)porphyrinato iron (III) complex (1.4 mmol) was dissolved in benzene (20 ml), and a small amount of 10% palladium black was added. A hydrogen gas was blown in for 20 minutes to reduce the central metal iron, and the catalyst was filtered off. A carbon monoxide gas was passed through this solution to convert the porphyrin iron complex to a carbon monoxide complex, and the solvent was distilled away under reduced pressure. The solid residue was dissolved in dimethyl sulfoxide solution (10 ml) with aeration of carbon monoxide gas, and human serum albumin in phosphate buffer (100 ml, pH 7.4, 1/30 mM) was added, which was followed by shaking. The mixture was concentrated to 20 ml by ultrafiltration (ultrafilter manufactured by ADVANTEC; cut off molecular weight 20,000). An aqueous solution of phosphate buffer (92 ml, pH 7.4, 1/30 mM) was added and the mixture was shaken, which was followed by concentration by ultrafiltration and dilution with an aqueous solution of phosphate buffer, whereby the desired dispersion of carbon monoxide-coordinated porphyrin iron (II) complex-albumin inclusion compound was obtained.

[0051] Irradiation of light (500 W) in an ice bath with nitrogen gas aeration gave a deoxy compound, whereby the objective dispersion of 2-(8'-(N-imidazolyl)octanoyloxymethyl)-5,10,15,20-tetrakis-(α,α,α,α-o-pivalamidophenyl)porphyrinato iron (II) complex-albumin inclusion compound was obtained.

[0052] This dispersion was diluted 1/50-fold and transferred into a quartz spectrometry cell, and the cell was sealed under a nitrogen atmosphere. The visible absorption spectra thereof were λ_{max} : 564 nm, 541 nm, 435 nm, which correspond to the spectrum of a penta-coordinated deoxy type complex having one intramolecular base.

[0053] The spectrum immediately changed when an oxygen gas was passed through this dispersion, giving spectra of λ_{max} : 546 nm, 423 nm, which apparently suggests that the porphyrin iron (II) complex included in albumin had changed to an oxygenated complex. A nitrogen gas was passed through the dispersion of this oxygenated complex, as a result of which the visible absorption spectrum changed from the oxygen type spectrum to the deoxy type spectrum, thus confirming the reversible adsorption and desorption of oxygen. Repeats of alternative aeration of oxygen and nitrogen enabled repetitive adsorption and desorption of oxygen. The half life time of the above-mentioned oxygenated complex was not less than 14 hours at 25°C.

Example 4

[0054] In the same manner as in Example 1 except that a recombinant human serum albumin (manufactured by The Green Cross Corporation) was used instead of human serum albumin, a dispersion of 2-(8'-(2"-methyl-1"-imidazolyl) octanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-pivalamidophenyl)porphyrinato iron (III) complex-recombinant human serum albumin inclusion compound was prepared. The visible absorption spectra were λ_{max} : 562 nm, 540 nm, 439 nm, which correspond to the spectrum of a deoxy type complex. The aeration of oxygen gas in the same manner as in Example 1 provided visible absorption spectra (λ_{max} : 545 nm, 423 nm) corresponding to the spectrum of an oxygenated complex. A nitrogen gas was passed through the dispersion of this oxygenated complex, whereby the original deoxy type spectrum was obtained, thus confirming the reversible adsorption and desorption of oxygen. The half life time of the above-mentioned oxygenated complex was not less than 14 hours at 25°C.

Example 5

[0055] 2-(8'-(N-Imidazolyl)octanoyloxymethyl)-5,10,15,20-tetrakis-(α,α,α,α-o-(2,2-dimethylhexanamido)phenyl) porphyrinato cobalt (II) complex (0.14 mmol) was dissolved in dimethyl sulfoxide (12 ml), and human serum albumin (0.1 mmol) in phosphate buffer (100 ml, pH 7.4, 1/30 mM) was added, which was followed by shaking. The mixture was concentrated to 20 ml by ultrafiltration (ultrafilter manufactured by ADVANTEC; cut off molecular weight 20,000). An aqueous solution of phosphate buffer (92 ml, pH 7.4, 1/30 mM) was added and the mixture was shaken, which was followed by concentration by ultrafiltration and dilution with an aqueous solution of phosphate buffer, whereby the

desired dispersion of porphyrin cobalt (II) complex-albumin inclusion compound was obtained. This dispersion was preserved at room temperature or 4°C for several months. As a result, no sedimentation or coagulation was found, and the dispersion was stable. The porphyrin cobalt (II) complex was converted to a deoxy type complex by the aeration of nitrogen gas. This dispersion was diluted 1/50-fold and transferred into a quartz spectrometry cell, and the cell was sealed under a nitrogen atmosphere. The visible absorption spectra thereof were λ_{max} : 530 nm, 411 nm, which correspond to the spectrum of a penta-coordinated deoxy type complex having one intramolecular base.

[0056] The spectrum immediately changed when an oxygen gas was passed through this dispersion, giving spectra of λ_{max} : 547 nm, 416 nm, which apparently suggests that the porphyrin cobalt (II) complex included in albumin had changed to an oxygenated complex. A nitrogen gas was passed through the dispersion of this oxygenated complex, as a result of which the visible absorption spectrum changed from the oxygen type spectrum to the deoxy type spectrum, thus confirming the reversible adsorption and desorption of oxygen. Repeats of alternative aeration of oxygen and nitrogen enabled repetitive adsorption and desorption of oxygen. The half life time of the above-mentioned oxygenated complex was not less than 11 hours at 25°C.

Example 6

[0057] 8,13-bis(1'-Methoxyethyl)-3,7,12,17-tetramethyl-2,18-bis(2'-methyloxycarbonylethyl)-21H,23H-porphyrin (0.34 mg, 0.52 mmol) synthesized by the method described in Tsuchida et al., Chem. Lett., 1953, (1994) was dissolved in 4N hydrochloric acid (4 ml), and the mixture was stirred for 20 minutes at room temperature. The reaction mixture was extracted with dichloromethane and washed with pure water. The organic layer was dried over anhydrous sodium sulfate, filtrated and concentrated to dryness under reduced pressure. The residue was separated and purified by silica gel column chromatography (dichloromethane/methanol=20/1 v/v) to give 165 mg of a monoester compound wherein one of the ester bonds was hydrolyzed (yield 50%).

[0058] This monoester compound (165 mg, 0.26 mmol) was dissolved in anhydrous tetrahydrofuran (5 ml) containing 4-dimethylaminopyridine (62.9 mg, 0.51 mmol), and the mixture was cooled to -15°C under a nitrogen atmosphere. Chloride pivalate (63 µl, 0.51 mmol) was added, and the mixture was stirred for 30 minutes. 1-(3-Aminopropyl)-imidazole (150 µl, 1.28 mmol) was added, and the mixture was reacted at -10 - 0°C for 1 hour and at room temperature for 4 hours. The solvent was removed under reduced pressure, and the residue was extracted with dichloromethane, which was followed by washing with pure water. Then, the organic layer was dried over anhydrous sodium sulfate, filtrated and concentrated to dryness under reduced pressure.

[0059] The residue was separated and purified by silica gel column chromatography (chloroform/methanol=20/1 v/v) to give 100 mg of 8,13-bis(1'-methoxyethyl)-2-(2'-methyloxycarbonylethyl)-18-(2'-(3"-imidazolyl)propyl)aminocarbonylethyl)-3,7,12,17-tetramethyl-21H,23H-porphyrin (yield 52%).

Elemental analysis (% by weight): C 69.4 (69.1), H 6.98 (7.14), N 13.4 (13.1) wherein the values in the parentheses are the calculated values of $C_{43}H_{53}N_7O_5$.

Thin-layer chromatography (Merck silica gel plate, chloroform/methanol=20/1 v/v): Rf: 0.42 (monospot)

FAB-mass spectrum: 748[M]$^+$

Infrared absorption spectrum (cm^{-1}): 1645 (ν_{c-o} (amide)), 1732 (ν_{c-o} (ester))

Visible absorption spectrum: (chloroform) λ_{max} : 623, 568, 533, 499, 402 nm

^{1}H NMR ($CDCl_3$, TMS standard), δ (ppm): 10.5, 10.0 (4H, m, meso-H), 6.9, 6.1 (3H, t, imidazole ring H), 6.0 (2H, s, $(CH_3O)C\underline{H}$-), 4.3 (4H, t, -$C\underline{H_2}CH_2COO$-), 3.8-3.6 (18H, m, CH_3O-, CH_3-), 3.3-3.1 (6H, t, -$CH_2C\underline{H_2}COO$-, -$CONHC\underline{H_2}$-), 2.9 (2H, t, -CH_2Im), 2.3 (6H, s, $(CH_3O)CHC\underline{H_2}$), 1.8 (3H, s, $COOCH_3$), 1.3 (2H, m, -$CH_2C\underline{H_2}C\underline{H_2}$-)

[0060] The porphyrin derivative (100 mg, 0.13 mmol) thus obtained was dissolved in dimethylformamide (10 ml). After thorough displacement with nitrogen gas, iron (II) chloride (266 mg, 1.34 mmol) was quickly added, and the mixture was reacted at 60°C for 2 hours. The solvent was removed under reduced pressure, and the obtained solid residue was dissolved in chloroform and washed with water. The organic layer was dehydrated over anhydrous sodium sulfate. The residue was filtered and the filtrate was concentrated to dryness under reduced pressure. The obtained solid residue was separated and purified by silica gel column chromatography (chloroform/methanol=5/1 v/v) to give the objective fractions, which were dried in vacuo, whereby 84 mg of the desired porphyrin iron complex was obtained (yield 78%).

Elemental analysis (% by weight): C 61.8 (61.7), H 6.41 (6.38), N 11.4 (11.7) wherein the values in the parentheses are the calculated values of $C_{43}H_{51}N_7O_5FeCl$.

Thin-layer chromatography (Merck silica gel plate, chloroform/methanol=10/1 v/v): Rf: 0.34 (monospot)

FAB-mass spectrum: 801[M-Cl]$^+$

Infrared absorption spectrum (cm^{-1}): 1652 (ν_{c-o} (amide)), 1728 (ν_{c-o} (ester))

Visible absorption spectrum: (chloroform) λ_{max}: 371, 500, 535 nm

[0061] To a solution of this complex (0.14 mmol) in dimethyl sulfoxide (12 ml) was added human serum albumin (0.1 mmol) in phosphate buffer (100 ml, pH 7.4, 1/30 mM), which was followed by shaking. The mixture was concentrated to 20 ml by ultrafiltration (ultrafilter manufactured by ADVANTEC; cut off molecular weight 20,000). An aqueous solution of phosphate buffer (92 ml, pH 7.4, 1/30 mM) was added and the mixture was shaken, which was followed by concentration by ultrafiltration and dilution with an aqueous solution of phosphate buffer, whereby the desired dispersion of porphyrin iron (III) complex-albumin inclusion compound was obtained. This dispersion was preserved at room temperature or 4°C for several months. As a result, no sedimentation or coagulation was found, and the dispersion was stable. A small amount of an aqueous solution of sodium dithionite was added to this dispersion under nitrogen atmosphere to reduce the central metal iron of the porphyrin complex to a divalent iron, whereby the objective dispersion of 8,13-bis(1'-methoxyethyl)-2-(2'-methoxycarbonylethyl)-18-(2'-(3"-imidazolyl)propyl)aminocarbonylethyl)-3,7,12,17-tetramethyl-porphyrinato iron (II) complex-albumin inclusion compound was obtained. This dispersion was diluted 1/50-fold and transferred into a quartz spectrometry cell, and the cell was sealed under a nitrogen atmosphere. The visible absorption spectra thereof were λ_{max} : 552 nm, 419 nm, which correspond to the spectrum of a penta-coordinated deoxy type complex having one intramolecular base.

[0062] The spectrum immediately changed when an oxygen gas was passed through this dispersion, giving spectra of λ_{max} : 567 nm, 536 nm, 408 nm, which apparently suggests that the porphyrin iron (II) complex included in albumin had changed to an oxygenated complex. The half life time of the above-mentioned oxygenated complex was within one hour at 25°C. A carbon monoxide gas was passed through the dispersion of the deoxy type complex, as a result of which the visible absorption spectra (λ_{max} : 560 nm, 631 nm, 412 nm) suggesting the formation of a carbon monoxide complex were obtained. This carbon monoxide complex was extremely stable at room temperature.

Example 7

[0063] In the same manner as in Example 5 except that 8,13-bis(1'-octadecanoxyethyl)-2-(2'-octadecanoxycarbonylethyl)-18-(2'-(10"-(2"'-methyl-1"'-imidazolyl)decane)aminocarbonylethyl)-3,7,12,17-tetramethyl-porphyrinato iron (III) complex was used instead of the porphyrin iron (III) complex used in Example 1 and bovine serum albumin was used instead of human serum albumin, a dispersion of 8,13-bis (1'-octadecanoxyethyl)-2-(2'-octadecanoxycarbonylethyl)-18-(2'-(10"-(2"'-methyl-1"'-imidazolyl)decane)aminocarbonylethyl)-3,7,12,17-tetramethyl-porphyrinato iron (II) complex-albumin inclusion compound was prepared. The visible absorption spectra were λ_{max} : 553 nm, 420 nm, which correspond to the spectrum of a deoxy type complex. The aeration of oxygen gas in the same manner as in Example 1 provided visible absorption spectra (λ_{max} : 567 nm, 536 nm, 407 nm) corresponding to the spectrum of an oxygenated complex. The half life time of the oxygenated complex was within one hour at 25°C. A carbon monoxide gas was passed through the dispersion of the deoxy type complex, whereby the spectra (λ_{max} : 560 nm, 531 nm, 413 nm) suggesting the formation of a carbon monoxide complex were obtained. This carbon monoxide complex was extremely stable at room temperature.

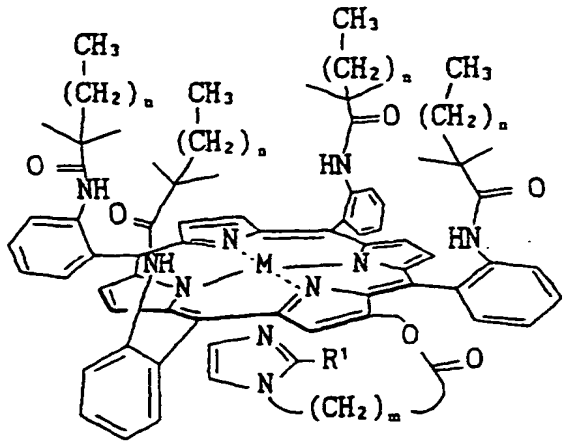
Experimental Example 1

[0064] The number of porphyrin iron complex bound to albumin in the inclusion compound of Example 1 as determined from the Scatchard plot was 1. The equilibrium constant was 2.6×10^6 (M^{-1}), thereby clarifying that the bond was almost as strong as in bilirubin and the like. The repetitive cycle of deoxy type complex and oxy type complex was observed more than 100 times at 25°C. The affinity for oxygen ($P_{1/2}$ (O_2)) was 24 Torr (37°C). The oxygen carrying efficiency between lung (110 Torr) and peripheral tissues (40 Torr) as estimated from the oxygen binding dissociation equilibrium curve was about 20% (37°C), thereby suggesting the effective action of the inclusion compound of the present invention as an oxygen carrier replacing erythrocytes. The behavior of the oxygen coordination complex in the inclusion compound of the present invention was clarified from the enthalpy/entropy changes caused by oxygen binding, and found to be about the same as hemoglobin in erythrocytes. The oxygen binding dissociation rate constant k_{on} and k_{off} as determined by laser flash photolysis was 2.4×10^8 ($M^{-1}s^{-1}$) and 3.2×10^3 (s^{-1}), respectively, thereby suggesting the ability of the inclusion compound of the present invention to bind or dissociate oxygen faster than erythrocytes. From the above results, it is clear that the inclusion compound of the present invention is capable of carrying oxygen sufficiently even when the compound is carried at high speeds in the body.

[0065] The substituted porphyrin metal complex-albumin inclusion compound of the present invention comprises a porphyrin metal complex as an oxygen adsorption/desorption site and functions as an oxygen carrier which is superior in biocompatibility, capable of adsorption and desorption of oxygen under physiological conditions, and is rather easily produced at an industrial scale.

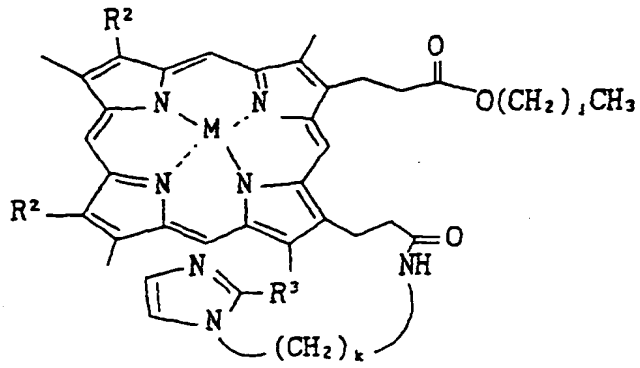
Claims

1. A porphyrin metal complex-albumin inclusion compound wherein a substituted porphyrin metal complex comprising, as a central, coordinated metal M an iron or cobalt ion, is included in the albumin,
wherein the substituted porphyrin metal complex is a 5,10,15,20-tetra[α,α,α,α-o-(substituted amide)phenyl]porphyrin metal complex of the following formula (I)



(I)

wherein R^1 is hydrogen or methyl, n is an integer of from 0 to 17 and m is an integer of from 3 to 17; or a porphyrin metal complex of the formula (II)



(II)

wherein the two R^2 groups are the same or different and each is a vinyl or a 1-alkyloxyethyl group of the formula : $-(CH_3)CHO(CH_2)_hCH_3$ wherein h is an integer of from 0 to 17, R^3 is hydrogen or methyl, j is an integer of from 0 to 17 and k is an integer of from 3 to 10.

2. The inclusion compound of Claim 1, wherein the iron or cobalt is divalent.

3. The inclusion compound of claim 1, wherein the substituted porphyrin metal complex is a 2-(8'-(2"-methyl-1"-imidazolyl)octanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-pivalamidophenyl)porphyrinato iron complex.

4. The inclusion compound of claim 1, wherein the substituted porphyrin metal complex is a 2-(18'-(N-imidazolyl) octadecanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-(2,2-dimethyloctadecanamidophenyl)porphyrinato iron complex, a 2-(8'-(N-imidazolyl)octanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-pivalamidophenyl)porphyrinato iron complex or a 2-(8'-(N-imidazolyl)octanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-(2,2-dimethylhexanami-

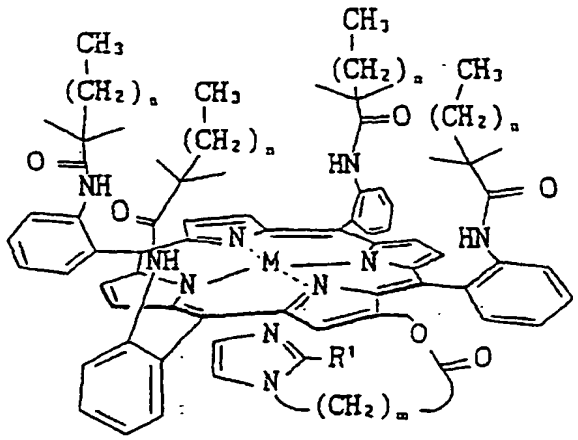
do)phenyl)porphyrinato cobalt complex.

5. The inclusion compound of any one of Claims 1 to 4, wherein the albumin is a human serum albumin or a recombinant human serum albumin.

6. An oxygen carrier composition comprising, as an active ingredient, a porphyrin metal complex-albumin inclusion compound according to any one of Claims 1 to 5.

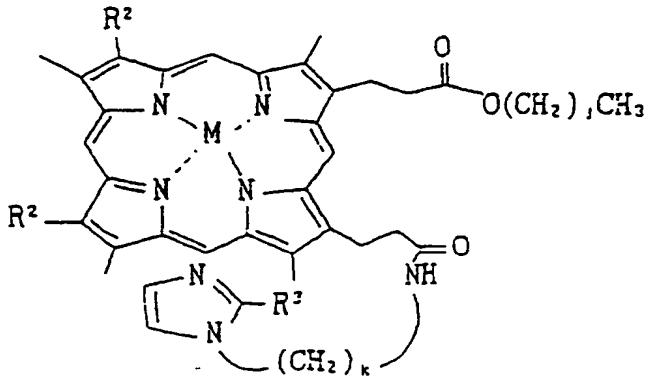
Patentansprüche

1. Porphyrinmetallkomplex-Albumin-Einschlussverbindung, in der ein substituierter Porphyrinmetallkomplex, der als zentrales koordiniertes Metall M ein. Eisen- oder Cobalt-Ion umfasst, im Albumin eingeschlossen ist, wobei es sich bei dem substituierten Porphyrinmetallkomplex um einen 5,10,15,20-Tetra[α,α,α,α-o-(substituiertes Amid) phenyl]porphyrinmetallkomplex der nachstehenden Formel (I) handelt



(I)

in der R^1 ein Wasserstoffatom oder eine Methylgruppe ist, n eine ganze Zahl von 0 bis 17 ist und m eine ganze Zahl von 3 bis 17 ist;
oder um einen Porphyrinmetallkomplex der Formel (II)



(II)

in der die beiden Reste R^2 gleich oder verschieden sind und jeder einen Vinylrest oder einen 1-Alkyloxyethylrest

der Formel $-(CH_3)CHO(CH_2)_hCH_3$ darstellt, wobei h eine ganze Zahl von 0 bis 17 ist, R^3 ein Wasserstoffatom oder ein Methylrest ist, j eine ganze Zahl von 0 bis 17 ist und k eine ganze Zahl von 3 bis 10 ist.

2. Einschlussverbindung nach Anspruch 1, wobei das Eisen oder Cobalt zweiwertig ist.

3. Einschlussverbindung nach Anspruch 1, wobei der substituierte Porphyrinmetallkomplex ein 2-(8'-(2"-Methyl-1"-imidazolyl)octanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-pivalamidophenyl)porphyrinato-Eisenkomplex ist.

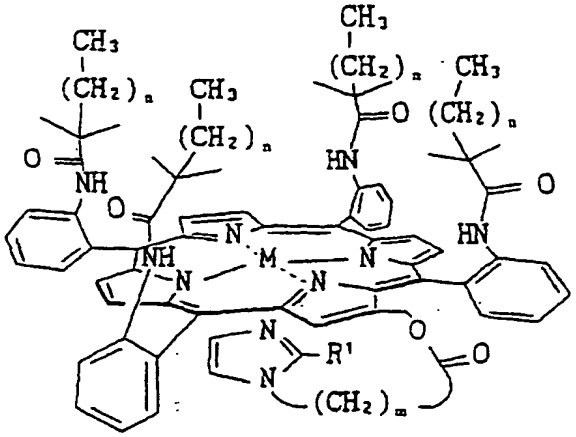
4. Einschlussverbindung nach Anspruch 1, wobei der substituierte Porphyrinmetallkomplex ein 2-(18'-(N-imidazolyl) octadecanoyloxymethyl)-5,10,15,20-tetrakis-(α,α,α,α-o-(2,2-dimethyloctadecanamidophenyl)porphyrinato-Eisenkomplex, ein 2-(8'-(N-imidazolyl)octanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-pivalamidophenyl)porphyrinato-Eisenkomplex oder ein 2-(8'-(N-imidazolyl)octanoyloxymethyl)-5,10,15,20-tetrakis(α,α,α,α-o-(2,2-dimethylhexanamido)phenyl)porphyrinato-Cobaltkomplex ist.

5. Einschlussverbindung nach einem der Ansprüche 1 bis 4, wobei das Albumin ein Human-Serumalbumin oder ein rekombinantes Human-Serumalbumin ist.

6. Sauerstoffträger-Zusammensetzung, die als wirksamen Bestandteil eine Porphyrinmetallkomplex-Albumin-Einschlussverbindung nach einem der Ansprüche 1 bis 5 umfasst.

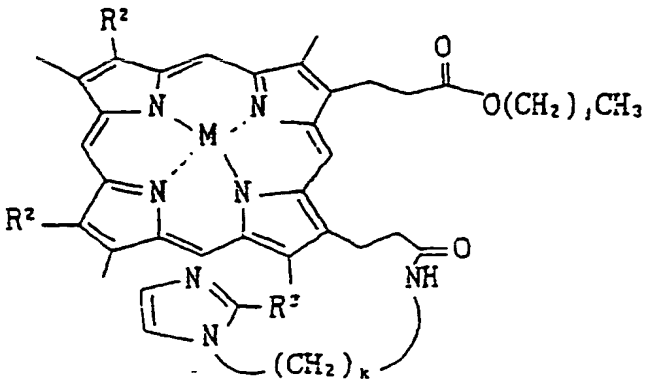
Revendications

1. Composé d'inclusion d'un complexe de porphyrine-métaux et de l'albumine dans lequel un complexe de porphyrine-métaux substitué comprenant, en tant que métal coordonné central M, un ion fer ou un ion cobalt, est inclus dans l'albumine, dans lequel le complexe de porphyrine-métaux substitué est un complexe de 5,10,15,20-tétra[α,α,α,α-o-(amide substitué)-phényl]porphyrine-métaux de formule (I) suivante :



(I)

dans laquelle R^1 est l'hydrogène ou un groupe méthyle, n est un entier valant de 0 à 17 et m est un entier valant de 3 à 17 ; ou
un complexe de porphyrine-métaux de formule (II) :



(II)

dans laquelle les deux groupes R^2 sont identiques eu différents et chacun est un groupe vinyle ou un groupe 1-alkyleoxyethyle de formule: $-(CH_3)CHO(CH_2)_hCH_3$, dans laquelle h est un entier valant de 0 à 17, R^3 est l'hydrogène ou un groupe méthyle, j est un entier valant de 0 à 17 et k est un entier valant de 3 à 10.

2. Composé d'inclusion selon la revendication 1, dans lequel le fer ou le cobalt est divalent.

3. Composé d'inclusion selon la revendication 1, dans lequel le complexe de porphyrine-métaux substitué est un complexe 2-(8'-(2"-méthyl-1"-imidazolyl)octanoyloxyméthyl)-5,10,15,20-tétrakis(α,α,α,α-o-pivalamidophényl) porphyrine-fer.

4. Composé d'inclusion selon la revendication 1, dans lequel le complexe de porphyrine-métaux substitué est un complexe de 2-(18'-(N-imidazolyl)octadécanoyloxyméthyl)-5,10,15,20-tétrakis (α,α,α,α-o-(2,2-diméthyloctadécanamidophényl)porphyrine-fer, un complexe de 2-(8'-(N-imidazolyl)octanoyloxyméthyl)-5,10,15,20-tétrakis(α,α,α,α-o-pivalamidophényl)porphyrine-fer ou un complexe de 2-(8'-(N-imidazolyl)octanoyloxyméthyl)-5,10,15,20-tétrakis(α,α,α,α-o-(2,2-diméthylhexanamido)phényl)porphyrine-cobalt.

5. Composé d'inclusion selon l'une quelconque des revendications 1 à 4, dans lequel l'albumine est une sérumalbumine humaine ou une sérumalbumine humaine recombinante.

6. Composition porteuse d'oxygène comprenant, en tant qu'ingrédient actif, un composé d'inclusion d'un complexe de porphyrine-métaux et de l'albumine selon l'une quelconque des revendications 1 à 5.

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(54) Porphyrin metal complex-albumin inclusion compound and oxygen carrier composition comprising the same

(57) A porphyrin metal complex-albumin inclusion compound wherein a substituted porphyrin metal complex comprising, as a central, coordinated metal, a transitional metal belonging to the fourth or fifth period of the periodic table, is included in the albumin, and an oxygen carrier composition comprising said inclusion compound as an active ingredient. The compound functions as an oxygen carrier which is superior in biocompatibility, capable of adsorption and desorption of oxygen under physiological conditions, and is rather easily produced at an industrial scale.

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EUROPEAN SEARCH REPORT

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DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
X,P	WO 96 03426 A (DUKE UNIVERSITY) 8 February 1996 * claims 1-20 *	1,5,6	A61K47/48 A61K9/14
X	BONAVENTURA J ET AL: "OXYGEN-BINDING ALBUMINS: A NOVEL APPROACH TO BLOOD SUBSTITUTES" ARTIFICIAL CELLS, BLOOD SUBSTITUTES, AND IMMOBILIZATION BIOTECHNOLOGY, vol. 22, no. 5, 1 January 1994, page A83 XP000576501 * the whole document *	1,5,6	
X	EP 0 601 183 A (MEIJI MILK PRODUCTS COMPANY LIMITED) 15 June 1994 * claims 1-9 * * figures 5,12 *	1,5,6	
A	PATENT ABSTRACTS OF JAPAN vol. 9, no. 12 (C-261), 18 January 1985 & JP 59 162924 A (H. TSUCHIDA), 13 September 1984 * abstract *	1-6	TECHNICAL FIELDS SEARCHED (Int.Cl.6) A61K
A	PATENT ABSTRACTS OF JAPAN vol. 18, no. 528 (C-1258), 6 October 1994 & JP 06 184156 A (RES INST FOR PROD DEV ET AL), 5 July 1994 * abstract *	1-6	
A,D	PATENT ABSTRACTS OF JAPAN vol. 18, no. 680 (C-1291), 21 December 1994 & JP 06 271577 A (RES INST OR PROD DEV), 27 September 1994 * abstract *	1-6	

-/--

The present search report has been drawn up for all claims

Place of search	Date of completion of the search	Examiner
BERLIN	29 January 1999	Siatou, E

CATEGORY OF CITED DOCUMENTS

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EUROPEAN SEARCH REPORT

Application Number
EP 96 10 6625

DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
A,D	J. P. COLLMAN ET AL: ""Picket Fence Porphyrins." Synthetic Models for Oxygen Binding Hemoproteins" JOURNAL OF THE AMERICAN CHEMICAL SOCIETY, vol. 97, January 1975 - March 1975, pages 1427-1439, XP002091695 * page 1429 *	1-6	
A,D	TRAYLOR T G ET AL: "SYNTHESES AND NMR CHARACTERIZATION OF CHELATED HEME MODELS OF HEMOPROTEINS" JOURNAL OF THE AMERICAN CHEMICAL SOCIETY, vol. 101, no. 22, 24 October 1979, pages 6716-6731, XP000575976 * page 6718; table I *	1-6	
			TECHNICAL FIELDS SEARCHED (Int.Cl.6)

The present search report has been drawn up for all claims

Place of search	Date of completion of the search	Examiner
BERLIN	29 January 1999	Siatou, E

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X : particularly relevant if taken alone
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This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
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29-01-1999

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9603426 A	08-02-1996	AU 685661 B	22-01-1998
		AU 3093795 A	22-02-1996
		CA 2195630 A	08-02-1996
		EP 0775157 A	28-05-1997
		JP 10503489 T	31-03-1998
		US 5773417 A	30-06-1998
EP 601183 A	15-06-1994	AU 663106 B	28-09-1995
		AU 2390392 A	02-03-1993
		CA 2113837 A	18-02-1993
		WO 9303035 A	18-02-1993
		US 5629198 A	13-05-1997

EPO FORM P0459

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